Under the Paperwork Reduction Act of 1995, no persons are required to

INFORMATION DISCLOSURE
STATEMENT BY APPLICANT
( Not for submission under 37 CFR 1.99)

Application Number		10790574
Filing Date		2004-03-01
First Named Inventor John		D. Mehr
Art Unit		2153
Examiner Name		
Attorney Docket Numb	er	MS307141.01/MSFTP573US

					U.S.	PATENTS			Remove	
Examiner Initial*	Cite No	Patent Number	Kind Code <sup>1</sup>	Issue E	Date	Name of Patentee or Applicant of cited Document			s,Columns,Lines where ant Passages or Releves Appear	
	1									
If you wis	h to a	dd additional U.S. Pater	nt citatio	n inform	ation pl	ease click the	Add button.		Add	
			U.S.P	ATENT	APPLI	CATION PUB	LICATIONS		Remove	
Examiner Initial*	Cite No	Publication Number	Kind Code <sup>1</sup>	Publica Date	ation	Name of Patentee or Applicant of cited Document  Bandini, et al.		Pages,Columns,Lines where Relevant Passages or Relevan Figures Appear		
	1	20020199095		2002-12	2-00					
	2	20030204569				Andrews, et a	L			
	3	20030009698				Lindeman, et al.				
If you wis	h to a	dd additional U.S. Publi	shed Ap	plication	citatio	n information p	lease click the Ad	d buttor		
				FOREIG	GN PAT	ENT DOCUM	ENTS		Remove	
Examiner Initial*	Cite No	Foreign Document Number <sup>3</sup>	Country Code <sup>2</sup>		Kind Code <sup>4</sup>	Publication Date Name of Pat Applicant of Document		e or	Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear	Ts
	1									

#### 

If you wish	h to a	dd addi1	tonal Foreign	Patent D	ocumer	nt citation	n informati	on please	e click the Add	button	Add		
				NO	N-PATI	ENT LIT	ERATUR	DOCUM	MENTS		Remove		
Examiner Initials*	Cite No	(book,		urnal, sei	rial, sym	posium,	catalog, e		article (when a , pages(s), vol				Тs
	1												
If you wis	h to a	dd addit	ional non-pate	ent literat	ture doc	ument c	itation info	rmation p	lease click the	Add b	utton Ad	id	
					E	XAMINE	R SIGNA	TURE					
Examiner	Signa	ture							Date Conside	ered			
*EYAMIN	ED: In	itial if re	forence cone	idered w	hathar	or not cit	ation is in	conform	ance with MDE	D RND	Draw line	through a	

1 See Kind Codes of USPTO Patent Documents at yeary\_USPTO\_GOV or MPEP 901.6.2 Extens office that issued the document, by the ho-other code (WIPD Standard ST.3.) 27% plaquese petited colouresh, by mandation of the year of the Preprior under process the sear injuries or fire period the Democrature process the sear injuries of the Patend document.

1 Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST.16 if possible. 2 Applicant is to place a check mark here if English language transition is attached.

citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

# INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		10790574
Filing Date		2004-03-01
First Named Inventor John		D. Mehr
Art Unit		2153
Examiner Name		
Attorney Docket Numb	er	MS307141.01/MSFTP573US

#### CERTIFICATION STATEMENT

Please see 37 CFR 1.97 and 1.98 to make the appropriate selection(s);

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. See 37 CFR 197(eVI).

ΩR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office is a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquiry, on learn of information contained in the information disclosure statement was known to garry individual designated in 37 CFR 1/5(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 1/3/(k).

See attached certification statement.

Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.

.7 None

### SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

_			
Signature	/Himanshu S. Amin/	Date (YYYY-MM-DD)	2006-07-31
Name/Print	Himanshu S. Amin	Registration Number	40894

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is for life and by the USPTO to process) an application. Confidentiality is governed by \$5 U.S. C.12 GA 37 CFR.

1.14. This collection is estimated to take I hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, u.S. Operatment of Commence, P. O. Box 1430, Alexandriu, V.S. 2213.1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, V.A. 2213.1450.

## Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) famishing of the information solicided is voluntary, and (3) the principal purpose for which the information is used by the U.S. Patient and Trademan Kollie is to process and/or examine your submission related to a patient application or patient. If you do not furnish the requested process and/or examine your submission related to a patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested to the patient process and/or examine your submission, which may related that the patient process and/or examine your submission, which may

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
  - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement neodiations.
  - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record partains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
  - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974. as amended, pursuant to 5 U.S.C. 552a(m.).
  - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
    may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
    to the Patent Cooperation Treaty.
  - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
  - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uting an inspection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 44 U.S.C. 2014 and 2016. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make relevant for the control of t
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S. C. 12(b) or issuance of a patient pursuant to 35 U.S. C. 15.1 Further, a record may be disclosed, subject to the limitations of 37 CFR.114, as a routine use, to the public if the record via flori of mapplication which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inseptions or an issued patient.
  - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.